

884

## EP1357977

Publication Title:

CHROMIUM CONTAINING COMPOSITIONS FOR THE TREATMENT OF DIABETES, THE REDUCTION OF BODY FAT, IMPROVEMENT OF INSULIN SENSITIVITY AND REDUCTION OF HYPERGLYCEMIA AND HYPERCHOLESTEREMIA

Abstract:

Abstract not available for EP1357977

Abstract of corresponding document: WO0224180

A composition for treating insulin-dependent diabetes, reducing body fat, improving insulin sensitivity, reducing hyperglycemia, and reducing hypercholesterolemia with at least one chromium complex and a conjugated fatty acid or conjugated fatty alcohol is disclosed. A method of treating a subject suffering from insulin-dependent diabetes by administering a composition that includes at least one chromium complex and a conjugated fatty acid or conjugated fatty alcohol is similarly provided. The administration of a composition containing an effective dose of at least one chromium complex and a conjugated fatty acid or conjugated fatty alcohol for the treatment of obesity is likewise provided.

Data supplied from the esp@cenet database - Worldwide d37

-----  
Courtesy of <http://v3.espacenet.com>

(19)



Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11)

**EP 1 357 977 B1**

(12)

**EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention  
of the grant of the patent:  
**21.07.2004 Bulletin 2004/30**

(51) Int Cl.7: **A61P 31/00, A61K 31/555,  
A61K 31/455, A61K 31/20,  
A61K 31/045**

(21) Application number: **01975262.5**

(86) International application number:  
**PCT/US2001/029422**

(22) Date of filing: **20.09.2001**

(87) International publication number:  
**WO 2002/024180 (28.03.2002 Gazette 2002/12)**

(54) **CHROMIUM CONTAINING COMPOSITIONS FOR THE TREATMENT OF DIABETES, THE  
REDUCTION OF BODY FAT, IMPROVEMENT OF INSULIN SENSITIVITY AND REDUCTION OF  
HYPERGLYCEMIA AND HYPERCHOLESTEREMIA**

CHROM ENTHALTENDE ZUSAMMENSETZUNG ZUR BEHANDLUNG VON DIABETES, ZUR  
VERBESSERUNG DER INSULINSENSITIVITÄT UND ZUR REDUKTION VON HYPERGLYKÄMIE,  
HYPERCHOLESTEROLÄMIE UND DES KÖRPERFETTANTEILS

COMPOSITIONS CONTENANT DU CHROME POUR LE TRAITEMENT DU DIABETE, LA  
REDUCTION DES GRAISSES CORPORELLES, L'AMELIORATION DE LA SENSIBILITE A  
L'INSULINE, LA REDUCTION DE L'HYPERGLYCEMIE ET LA REDUCTION DE  
L'HYPERCHOLESTEROLEMIE.

(84) Designated Contracting States:  
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU  
MC NL PT SE TR**

• **KOMOROWSKI, James, R.**  
Trumbull, CT 06611 (US)  
• **GREENBERG, Danielle**  
Waccabuc, NY 10597 (US)

(30) Priority: **21.09.2000 US 234474 P**  
**06.06.2001 US 296688 P**

(74) Representative: **Tanner, James Percival et al**  
**D. Young & Co,**  
**21 New Fetter Lane**  
**London EC4A 1DA (GB)**

(43) Date of publication of application:  
**05.11.2003 Bulletin 2003/45**

(73) Proprietor: **Nutrition 21, Inc.**  
**Purchase, NY 10577-2197 (US)**

(56) References cited:  
**WO-A-00/12080 WO-A-01/43758**  
**WO-A-96/23811 WO-A-98/33494**  
**US-A- 5 626 849**

(72) Inventors:  
• **KATZ, David, P.**  
**Dobbs Ferry, NY 10522 (US)**

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

**EP 1 357 977 B1**

**Description**Background of the Invention5 Field of the Invention

[0001] This application claims priority to provisional application filed September 21, 2000 having application number US 60/234,474, and provisional application filed June 6, 2001 having application number US 60/296,688. The disclosed invention related to compositions useful for the treatment of type 1 diabetes, reduction of body fat, improvement of  
10 insulin sensitivity, reduction of hyperglycemia, and reduction of hypercholesterolemia. Specifically, compositions comprising chromium complexes in combination with conjugated compounds such as isomers of conjugated fatty acids or conjugated fatty alcohols.

Description of the Related Art15 Insulin-Dependent Diabetes

[0002] Diabetes is a chronic metabolic disorder which afflicts 16 million people in the United States, over one and one half million of whom have its most severe form, childhood diabetes (also called juvenile, type 1 or insulin-dependent  
20 diabetes). Insulin-dependent diabetes appears suddenly, most often in children and young adults, and progresses rapidly. In this form, the pancreas ceases to manufacture insulin, a hormone necessary to convert the food we eat into energy for the body. In the United States, diabetes is the fourth leading cause of death, killing more than 162,000 people each year. Notably, the mortality rate of patients with insulin-dependent diabetes increases dramatically after 15 years of disease duration. In addition, virtually every major organ system in the body is damaged by diabetes.  
25 Complications can include blindness, kidney failure, heart disease, stroke, amputation of extremities, loss of nerve sensation, early loss of teeth, high-risk pregnancies and babies born with birth defects.

[0003] Insulin resistance is characterized by reductions of glucose uptake in skeletal muscle. Currently, insulin injection is the only treatment method available for the over 1.5 million type 1 diabetics and becomes the eventual course of treatment for many of the more than 16 million type 2 diabetics in the United States. Nutritional therapies that positively  
30 impact glucose uptake in the face of insulin insufficiency would have a major impact on the long term treatment costs associated with diabetic care.

Obesity

[0004] More than half of U.S. adults are overweight and nearly one-quarter of the U.S. adults are considered to be obese. The increasing prevalence of overweight and obesity is a major public health concern, since obesity is associated with several chronic diseases. For example, overweight and obesity are known risk factors for diabetes, heart  
35 disease, stroke, hypertension, gallbladder disease, osteoarthritis, sleep apnea, and some forms of cancer such as uterine, breast, colorectal, kidney, and gallbladder. Furthermore, obesity is associated with high cholesterol, complications of pregnancy, menstrual irregularities, hirsutism, and increased surgical risk.  
40

[0005] Drugs currently approved by the FDA for the treatment of obesity produce weight losses of about 10% of initial body weight at one year when used singly. Combination therapy with phentermine and fenfluramine produced weight losses of about 15% of initial body weight at one year. Phenylpropanolamine (PPA) is an over-the-counter drug that has not been tested for long term use and is recommended for use for only about 12 weeks. With the exception  
45 of PPA, all of these drugs require a physician's prescription and are generally quite expensive. Side effects occur with all these drugs. For example, the administration of fenfluramine and phentermine for the treatment of obesity resulted in cardiac valve damage in some patients and ultimately led to the withdrawal of fenfluramine from the market. Two of the newest drugs for the treatment of obesity have side effects that limit their use. Sibutramine increases blood pressure in a subset of patients, and orlistat may have unpleasant gastrointestinal side effects.  
50

The Role of Chromium Complexes in the Treatment of Insulin-Dependent Maladies

[0006] Chromium picolinate is reported to produce modest weight loss and changes in body composition (Kaats, 1998, Cefalu, 1999). Chromium is a nutritionally essential trace element. The essentiality of chromium in the diet was  
55 established in 1959 by Schwartz, as cited in *Present Knowledge in Nutrition*, page 571, fifth edition (1984, the Nutrition Foundation, Washington, DC). Chromium depletion is characterized by the disturbance of glucose, lipid and protein metabolism and by a shortened lifespan. Chromium is essential for optimal insulin activity in all known insulin-dependent systems (Boyle et al., *Southern Med. J.* 70:1449-1453, 1977). Insufficient dietary chromium has been linked to both

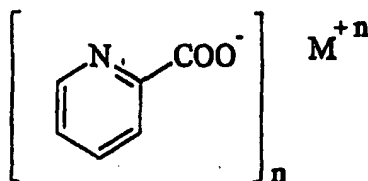
maturity-onset diabetes and to cardiovascular disease.

[0007] The principal energy sources for the body are glucose and fatty acids. Chromium depletion results in biologically ineffective insulin and compromised glucose metabolism. Under these conditions, the body must rely primarily on lipid metabolism to meet its energy requirements, resulting in the production of excessive amounts of acetyl-CoA and ketone bodies. Some of the documented acetyl-CoA is diverted to increased cholesterol biosynthesis, resulting in hypercholesterolemia. Diabetes mellitus is characterized in large part by glycosuria, hypercholesterolemia, and often ketoacidosis. The accelerated atherosclerotic process seen in diabetics is associated with hypercholesterolemia (Boyle et al., *supra*).

[0008] Dietary supplementation of chromium to normal individuals has been reported to lead to improvements in glucose tolerance, serum lipid concentrations, including high-density lipoprotein cholesterol, insulin and insulin binding (Anderson, *Clin. Psychol. Biochem.* 4:31-41, 1986). Supplemental chromium in the trivalent form, e.g. chromic chloride, is associated with improvements of risk factors associated with adult-onset (type 2) diabetes and cardiovascular disease.

[0009] Chromium functions as a cofactor for insulin. It binds to the insulin receptor and potentiates many, and perhaps all, of its functions (Boyle et al., *supra*). These functions include, but are not limited to, the regulation of carbohydrate and lipid metabolism. (*Present Knowledge in Nutrition, supra*, at p. 573-577). The introduction of inorganic chromium compounds *per se* into individuals is not particularly beneficial. Chromium must be converted endogenously into an organic complex or must be consumed as a biologically active molecule. Only about 0.5% of ingested inorganic chromium is assimilated into the body (*Recommended Daily Allowances*, Ninth Revised Edition, The National Academy of Sciences, page 160, 1980). Only 1-2% of most organic chromium compounds are assimilated into the body.

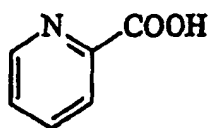
[0010] U.S. Patent No. Re. 33,988 discloses that when selected essential metals, including chromium, are administered to mammals as exogenously synthesized coordination complexes of picolinic acid, they are directly available for absorption without competition from other metals. This patent describes a composition and method for selectively supplementing the essential metals in the human diet and for facilitating absorption of these metals by intestinal cells. These complexes are safe, inexpensive, biocompatible and easy to produce. These exogenously synthesized essential metal coordination complexes of picolinic acid (pyridine-2-carboxylic acid) have the following structural formula:



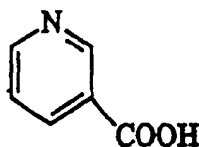
wherein M represents the metallic cation and n is equal to the cation's valence. For example, when M is Cr and n=3, then the compound is chromic tripicolinate. Other chromium picolinate complexes disclosed include chromic monopicolinate and chromic dipicolinate.

[0011] The U.S. Recommended Daily Intake (RDI) of chromium is 120 µg. U.S. Patent No. 5,087,623, describes the administration of chromic tripicolinate for the treatment of adult-onset diabetes in doses ranging from 50 to 500 µg. International Patent Application No. WO96/35421 discloses the use of high doses of chromic tripicolinate (providing 1,000-10,000 µg chromium/day) for reducing hyperglycemia and stabilizing the level of serum glucose in humans with type 2 diabetes. U.S. Patent No. 5,789,401 discloses a chromic tripicolinate-biotin composition and its use in lowering blood glucose levels in humans with type 2 diabetes.

[0012] U.S. Patent Nos. 5,087,623; 5,087,624; and 5,175,156, disclose the use of chromium tripicolinate for supplementing dietary chromium, reducing hyperglycemia and stabilizing serum glucose, increasing lean body mass and reducing body fat, and controlling blood serum lipid levels, including the lowering of undesirably high blood serum LDL-cholesterol levels and the raising of blood serum HDL-cholesterol levels. U.S. Patent Nos. 4,954,492 and 5,194,615, describe a related complex, chromic nicotinate, which is also used for supplementing dietary chromium and lowering serum lipid levels. Picolinic acid and nicotinic acid are position isomers having the following structures:



picolinic acid



nicotinic acid

[0013] Nicotinic acid and picolinic acid form coordination complexes with monovalent, divalent and trivalent metal ions and facilitate the absorption of these metals by transporting them across intestinal cells and into the bloodstream. Chromium absorption in rats following oral administration of  $\text{CrCl}_3$  was facilitated by the non-steroidal anti-inflammatory drugs (NSAIDs) aspirin and indomethacin (Davis et al., *J. Nutrition Res.* 15:202-210, 1995; Kamath et al., *J. Nutrition* 127:478-482, 1997). These drugs inhibit the enzyme cyclooxygenase which converts arachidonic acid to various prostaglandins, resulting in inhibition of intestinal mucus formation and lowering of intestinal pH which facilitates chromium absorption.

[0014] Additional pharmacological treatments for disorders caused by or exacerbated by improper glucose uptake are needed. Specifically, compositions for the treatment of diabetes and obesity would be a great boon to subjects suffering from these disease states. A new, more effective, less expensive treatment for diabetes and obesity with minimal side effects would be a great benefit to the treatment and prevention of obesity.

#### Brief Description of the Figures

[0015]

Figure 1 shows the effect of conjugated linoleic acid (CLA) on glucose uptake in human skeletal muscle culture.

Figure 2 graphs the effect of CP +CLA on glucose uptake in human skeletal muscle culture.

Figure 3 illustrates the effect of CP + CLA on glucose uptake in human skeletal muscle culture.

#### Summary of the Invention

[0016] The disclosed invention is directed to compositions useful for treating insulin-dependent diabetes, non-insulin dependent diabetes, reduction of body fat, improvement of insulin sensitivity, reduction of hyperglycemia, and reduction of hypercholesterolemia. The disclosed invention is a composition that includes at least one chromium complex and a conjugated fatty acid or conjugated fatty alcohol. Preferably, the chromium complex is chromium picolinate or chromium nicotinate. Advantageously, the conjugated fatty acid is conjugated linoleic acid.

[0017] In one aspect of the invention, the composition optionally includes a chelating agent. The chelating agent may be picolinic acid, nicotinic acid, or both.

[0018] The recited compositions are incorporated into a pharmaceutically effective carrier. The pharmaceutically effective carrier may be a tablet, capsule, microbead, emulsion, powder, granule, suspension, syrup or elixir. Preferably, when the carrier is a microbead, the microbead is a sugar beadlet or microcrystalline cellulose beadlet. Advantageously, the chromium complex and conjugated linoleic acid are coated on the beadlet. Optionally, the tablet, capsule, or microbead is coated with an enteric coating.

[0019] In another aspect of the invention, the chromium complex and conjugated fatty acid or conjugated fatty alcohol are in a ratio of between about 1:666 to about 1:200 (w/w).

[0020] Additional components may be included in the composition. For example, the composition may include at least one of a cyclooxygenase inhibitor, a mucolytic, or a salicin-containing herb. The cyclooxygenase inhibitor may be indomethacin, ibuprofen, acetaminophen, or naproxen. Advantageously, the salicin-containing herb may include *Boswellia serrata* (frankincense), *Betula lenta* (sweet birch), *Betula pubescens* (white birch), *Filipendula ulmaria* (meadowsweet), *Gaetheria procumbens* (wintergreens), *Polulus balsamifera*, *Populus jackii* (balm of Gilead) and *Salix alba* (white willow). The mucolytic may be guaifenesin.

[0021] Embodiments of the invention can be used in a method of treating type 1 diabetes in a subject. The method of treatment includes administering to a subject a pharmaceutically effective dose of a conjugated fatty acid or conjugated fatty alcohol in conjunction with at least one chromium complex. The chromium complex may be chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, or chromium yeasts. Optionally, the composition further includes at least one chelating agent. The chelating agent may be picolinic acid, nicotinic acid, or both.

[0022] A use for treating obesity in a subject is similarly contemplated. The method of treatment includes administering to a subject a pharmaceutical effective dose of conjugated fatty acid or conjugated fatty alcohol in conjunction with at least one chromium complex. The chromium complex may be chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, or chromium yeasts. Optionally, the composition further includes at least one chelating agent. The chelating agent may be picolinic acid, nicotinic acid, or both.

#### Detailed Description of the Preferred Embodiment

[0023] The disclosed invention relates to compositions for the treatment of type 1 diabetes. In particular, these compositions can be administered to reduce or even eliminate the need for administration of insulin in certain patients with type 1 diabetes. Further, they are used in methods for facilitating glucose uptake and utilization by cells. Certain individuals with type 2 diabetes will require insulin. The invention also relates to the treatment of type 2 diabetics who require insulin. Additionally, methods for treating obesity and increasing lean body mass are likewise contemplated. A primary basis of the present invention is the novel and unexpected discovery that compositions comprising an effective dose of a chromium complex in combination with a conjugated fatty acid or conjugated fatty alcohol produce a synergistic effect on glucose uptake in a subject in need thereof. Additionally, the co-administration of a chromium complex with a conjugated fatty acid or conjugated fatty alcohol, or an isomer thereof, demonstrate a synergistic, weight loss effect.

[0024] The terminology used in the description presented herein is not intended to be interpreted in any limited or restrictive manner, simply because it is being utilized in conjunction with a detailed description of certain specific embodiments of the invention. Furthermore, embodiments of the invention may include several novel features, no single one of which is solely responsible for its desirable attributes or which is essential to practicing the invention herein described.

[0025] The term "conjugated compound" refers to a compound having at least a portion that is a hydrocarbon, with at least three consecutive carbon-carbon bonds, such that single and double carbon-carbon bonds are found in an alternating manner. Thus, the compound will include the subunit  $C=C-C=C$  (hydrogens and other substituents not shown). Two preferred categories of conjugated compounds are fatty acids and fatty alcohols. It should be noted that these di- or poly-unsaturated compounds are referred to herein using the common names of the corresponding naturally-occurring compounds having the same number of carbons and unsaturations. Although such naturally-occurring compounds are not necessarily conjugated, due to the arrangement of their carbon-carbon double bonds, it will be understood in the context of the present invention that only conjugated versions of those compounds are contemplated; i.e., the arrangement of the double bonds will be such that they contain the substructure  $C=C-C=C$ . While compounds having as few as 4, 5, 6, or 7 carbon atoms are contemplated, the preferred conjugated compounds have 8, 9, 10, 12, 14, 16 or more carbon atoms, preferably not more than 32, 30, 28, or 26 carbon atoms. It should be noted that the phrase "conjugated fatty acid" or "conjugated fatty alcohol", as used herein, also includes isomers of fatty acids and fatty alcohols, as well as any other polyunsaturated compounds which act synergistically with chromium complexes to promote glucose uptake in a subject in need thereof. Suitable conjugated fatty acids include, without limitation, conjugated versions of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosatetraenoic acid, 9,12-octadecadienoic acid, octadecatrienoic acid, eicosatetraenoic acid, elcosapentaenoic acid, docosahexaenoic acid, docosapentaenoic acid, and all other diunsaturated and polyunsaturated fatty acids. In a preferred embodiment, the conjugated fatty acid is conjugated linoleic acid (CLA). As used herein, the phrase "conjugated fatty alcohols" includes, without limitation, conjugated versions of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, docosapentaenoic alcohol, and all other diunsaturated and polyunsaturated fatty alcohols. The present invention further includes the use of other conjugated compounds having at least 4, 5, 6, 7, or 8 carbon atoms, that functions synergistically with a chromium complex to facilitate glucose uptake and usage by cells. (Such function can be readily assayed, as further described below.) Note that the present invention includes alcohols and acids in which one or more of the double bonds result in a cis isomer, as well as those in which one or more of the double bonds result in a trans isomer. In some cases, all the double bonds are cis, while in others they are all trans, and in still other cases they are mixed cis and trans compounds.

[0026] In a preferred embodiment, the conjugated fatty acid of the composition is conjugated linoleic acid. Conjugated linoleic acid (CLA) has been reported in numerous publications to reduce fat gain in growing animals by 25%-70% (Park et al., 1997). Minimal studies have been performed on human subjects. In a preliminary study by Atkinson et al., there was no effect of CLA alone in obese subjects when added to standard diet and exercise. However, a subgroup of subjects who did not follow the diet and exercise program well, and who gained lean body mass, had a reduction in fat mass in the CLA subjects and an increase in fat mass in placebo subjects (Atkinson, communication, 2000). These

data, along with the animal data, suggest that CLA is most effective in preventing fat accumulation rather than producing fat loss. Two hypotheses have been advanced to explain the role of CLA in weight loss: 1) CLA may act as a  $\beta$ -3 agonist; and 2) CLA alters nutrient partitioning within muscles to promote fat oxidation rather than carbohydrate (CHO) oxidation (Park et al., 1999).

[0027] Embodiments of the present invention include compositions which contain a conjugated fatty acid or conjugated fatty alcohol in combination with at least one chromium complex. Additionally, uses of the compositions disclosed herein to treat diabetes and/or obesity are provided. The co-administration of a conjugated fatty acid or conjugated fatty alcohol and at least one chromium complex provides a number of advantages over conventional pharmaceutical regimes for the treatment of diabetes and promotion of weight loss. One advantage of the compositions described herein over conventional treatment regimes for diabetes (such as injection of insulin) and weight loss is that compositions comprising a conjugated compound such as a conjugated fatty acid or conjugated fatty alcohol and at least one chromium complex have no known side effects while still providing efficacy in glucose uptake and weight loss. With reference to Figure 1, one will observe that a conjugated fatty acid such as CLA alone had no effect on glucose uptake in human skeletal muscle culture under both basal (without insulin) and stimulated (with insulin) conditions. The units referred to on the vertical axis of Figures 1 and 2 are pico Moles per milligram of glucose uptake. The unit referred to on the vertical axis of Figure 3 is the percentage increase in the uptake of glucose in human skeletal muscle culture. The term Basal as used in Figures 1 and 2 refers to human skeletal muscle culture without insulin. Whereas the term Stimulated indicates human skeletal muscle culture with insulin. Those bars on Figures, 1, 2, and 3 representing Chromium Picolinate (CP) and Conjugated Linoleic Acid (CLA) are measured in nanograms per milliliter.

[0028] However, the administration of CLA in combination with a chromium complex such as chromium picolinate (CP) (as depicted in Figure 2) significantly increases glucose uptake in human skeletal muscle culture under both basal (without insulin) and stimulated (with insulin) conditions. CP and CLA under basal conditions was shown to have a greater effect than insulin alone.

[0029] Finally, turning to Figure 3, it will be appreciated that CLA plus a chromium complex synergistically aid in the percent increase in the uptake of glucose in human skeletal muscle culture.

[0030] In some embodiments, compositions comprising an elective dose of a chromium complex are provided. As used herein, the terms "chromium complexes" or "chromium complex" include, without limitation, chromium picolinate, chromic tripicolinate, chromium nicotinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts. Chromium picolinate, for example, has been shown to produce modest weight loss and changes in body composition.

[0031] The compositions of the disclosed invention additionally include an effective dose of conjugated compound such as a conjugated fatty acid or conjugated fatty alcohol, of the type previously described. In preferred embodiments, the conjugated fatty acid component of the composition is linoleic acid (CLA). Suitable forms of CLA include isomers with double bonds at carbons 9, 11; carbons 10,12; or a mixture thereof. Preferably, the compositions include the 10,12 form of CLA to reduce body fat and increase lean body mass. Advantageously, the composition will include the conjugated compound at a dose of at least 50% by weight of formulation. More preferably, the effective dose of the conjugated compound for a 70 kg human is between about 100 mg to about 10 grams. In a most preferred embodiment, the effective dose of the conjugated compound is 75 mg, 100 mg, 250 mg, 500 mg, 750 mg, 1000 mg, 2 g, 5 g, 7 g, or 10 g.

[0032] While the chromium complexes aid in the absorption of chromium by intestinal cells, in some embodiments, chelating agents are advantageously included in the compositions to facilitate absorption of other ingested chromium as well as other metals including, but not limited to, copper, iron, magnesium, manganese and zinc. Suitable chelating agents include picolinic acid, nicotinic acid, or both picolinic acid and nicotinic acid. Thus, the compositions of the disclosed invention are readily absorbable forms of chromium which also facilitate absorption of other essential metals in the human diet.

[0033] The chromium complexes of the disclosed invention have the same uses as described for chromic tripicolinate in U.S. Patent Nos. 5,087,623, 5,087,624 and 5,174,156, namely supplementing dietary chromium, lowering blood glucose levels in diabetics, lowering serum lipid levels and increasing lean body mass. Additionally, the chromium complexes of the present invention act to treat symptoms associated with overweight and obesity.

[0034] Advantageously, the chromium complexes are synthetic. The synthesis and use of chromium picolinate is described in U.S. Patent Nos. Re. 33,988 and 5,087,623. Chromic tripicolinate is available from health food stores, drug stores and other commercial sources. The synthesis and use of chromic polynicotinate is described in U.S. Patent No. 5,194,615.

[0035] The chelating agents such as picolinic acid and nicotinic acid are available from many commercial sources, including Sigma-Aldrich (St. Louis, MO) (picolinic acid; catalog No. P5503; nicotinic acid; catalog No. PN4126). Preferably, the ratio of the chromium complex to the chelating agent from about 10:1 to about 1:10 (w/w), more preferably from about 5:1 to about 1:5 (w/w).

[0036] A variety of delivery systems are available to deliver the compositions to a subject in need thereof. Preferably,

the compositions of the disclosed invention are prepared by incorporating the components into a pharmaceutically acceptable carrier, including but not limited to tablets, capsules and microbeads, preferably sugar beadlets or microcrystalline cellulose.

**[0037]** For oral administration, the chromium complex may be incorporated into a tablet, aqueous or oil suspension, dispersible powder or granule, microbead, emulsion, hard or soft capsule, syrup or elixir. The components of the composition may also be administered separately. Compositions may be prepared according to any method known in the art for the manufacture of pharmaceutically acceptable compositions and such compositions may contain one or more of the following agents: sweeteners, flavoring agents, coloring agents and preservatives. Tablets containing the active ingredients in admixture with nontoxic pharmaceutically acceptable excipients suitable for tablet manufacture are acceptable. "Pharmaceutically acceptable" means that the agent should be acceptable in the sense of being compatible with the other ingredients of the formulation (as well as non-injurious to the individual). Such excipients include inert diluents such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, such as corn starch and alginic acid; binding agents such as starch, gelatin or acacia; and lubricating agents such as magnesium stearate, stearic acid or talc. Tablets may be uncoated or may be coated with known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period of time. For example, a time delay material such as glyceryl monostearate or glyceryl stearate alone or with a wax may be employed.

**[0038]** In another preferred embodiment, tablets, capsules or microbeads are coated with an enteric coating which prevents dissolution in the acidic environment of the stomach. Instead, this coating dissolves in the small intestine at a more neutral pH. Because certain chromium complexes may be more stable at this neutral pH than at the acidic pH of the stomach, enhanced absorption occurs because the chromium complexes remain substantially intact until they reach the small intestine. Such enteric coated compositions are described by Bauer et al., *Coated Pharmaceutical Dosage Forms: Fundamentals, Manufacturing Techniques, Biopharmaceutical Aspects, Test Methods and Raw Materials*, CRC Press, Washington, DC, 1998.

**[0039]** Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, such as peanut oil, liquid paraffin or olive oil.

**[0040]** Aqueous suspensions may contain the chromium complexes of the invention in admixture with excipients for the manufacture of aqueous suspensions. Such excipients include suspending agents, dispersing or wetting agents, one or more preservatives, one or more coloring agents, one or more flavoring agents and one or more sweetening agents such as sucrose or saccharin.

**[0041]** Oil suspensions may be formulated by suspending the active ingredient in a vegetable oil, such as arachis oil, olive oil, sesame oil or coconut oil, or in a mineral oil such as liquid paraffin. The oil suspension may contain a thickening agent, such as beeswax, hard paraffin or cetyl alcohol. Sweetening agent, such as those set forth above, and flavoring agents may be added to provide a palatable oral preparation. These compositions may be preserved by an added antioxidant such as ascorbic acid. Dispersible powders and granules of the invention suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, a suspending agent, and one or more preservatives. Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

**[0042]** Syrups and elixirs may be formulated with sweetening agents, such as glycerol, sorbitol or sucrose. Such formulations may also contain a demulcent, a preservative, a flavoring or a coloring agent.

**[0043]** The oral formulations described above may also include aspirin (acetylsalicylic acid), other salicylates, or another NSAID such as indomethacin, ibuprofen, acetaminophen, naproxen or any drug capable of inhibiting the cyclooxygenase pathway leading to prostaglandin synthesis. This results in a decrease in intestinal mucus production and lower intestinal pH which facilitates absorption of the chromium compositions of the present invention. The oral compositions may further include mucolytics such as guaifenesin and the like, to inhibit intestinal mucus production, and/or acids such as ascorbic acid, citric acid and the like to lower intestinal pH. Inclusion of one or both of these compounds further enhances chromium absorption. There are two forms of cyclooxygenase (cox), cox1 and cox2, which differ in their sensitivity to inhibition by NSAIDs. The cox2 isozyme promotes prostaglandin formation at sites of inflammation, but not at other sites such as the gastrointestinal tract. In contrast, relatively selective inhibition of cox1 facilitates chronic tripicolinate and chronic polynicotinate absorption. Although the selective inhibition of cox1 is desirable, any inhibitor of cox1 or cox2 can be formulated with the chronic tripicolinate and chronic polynicotinate compositions of the invention. Cox inhibitors, acids and mucolytics may also be coadministered with the chronic tripicolinate and chronic polynicotinate compositions of the invention. The amount of these drugs formulated with or coadministered with the chronic tripicolinate compositions of the invention are as follows: cox inhibitions, between about 50 mg and 500 mg; mucolytics, between about 10 mg and 250 mg; and acids, between about 50 mg and about 1,000 mg.

**[0044]** The coadministration or formulation of salicylate-containing herbs with the compositions of the invention is also contemplated. Class I herbs, as documented in the American Herbal Products Association's *Botanical Safety*



*Handbook* (herbs that can be safely consumed when used appropriately), such as *Boswellia serrata* (frankincense), *Betula lenta* (sweet birch), *Betula pubescens* (white birch), *Filipendula ulmaria* (meadowsweet), *Gaultheria procumbens* (wintergreens), *Populus balsamifera* and *Populus jackii* (balm of Gilead), and *Salix alba* (white willow) are all salicin-containing plants with salicylate-like properties. These herbs suppress prostaglandin synthesis by cox inhibition, thereby improving absorption of the chromium complexes of the invention. These herbs are relatively free from gastric ulcerogenic effects (Singh et al., *Agents and Actions* 18:407-412, 1986). In addition, pre-clinical acute toxicity studies have shown that salicin-containing plants do not cause hematological disturbances (American Herbal Products Association, *Botanical Safety Handbook*, 1997).

[0045] The compounds and herbs described above all effect gut physiology by inhibiting prostaglandin synthesis, decreasing mucus production, and lowering gastrointestinal pH. The inclusion of these compounds, as well as an enteric coating, into the oral chromium picolinate or chromium nicotinate compositions of the invention results in a multicomponent delivery system which allows delivery of these agents to the gastrointestinal tract where they work in concert to facilitate chromium absorption.

[0046] In a preferred embodiment, the chromium complex is coated onto microbeads. in a particularly preferred embodiment, these microbeads are sugar beadlets of various sizes, also known as nonpareils, and are commercially available from, for example, SmithKline Beecham. If the microbeads are to be used to administer the compositions of the invention to diabetic patients, the administration of other types of microbeads, such as microcrystalline cellulose, is preferred. Microcrystalline cellulose is commercially available and can be processed into beadlets of various sizes by micronization, a technique well known in the art. The microbeads are essentially a carrier for the compositions of the invention. For a description of coated beadlets, see, for example, Carstensen, J. T., *Pharmaceutical Principles of Solid Dosage Forms*, Technomic Publishing Co., Inc., Lancaster, PA, pp. 228-230, 1993. Aqueous solutions containing the chromium complexes with or without the chelating agent components such as nicotinic acid and picolinic acid are sprayed onto the microbeads by well known methods. by suspending the microbeads in an upcurrent of air and introducing a fine spray of the active ingredients which form a coating on the outside of the microbeads which is then allowed to dry. The desired chromium complex components with or without a chelating agent may be combined into one same solution or applied using separate solutions. Optionally, the coated microbeads can be further coated with a substance to protect the active ingredients coated onto the beads, such as latex. The microbeads may be placed in a capsule prior to administration. In another preferred embodiment, the capsule or the microbeads are coated with an enteric coating to delay dissolution until reaching the small intestine.

[0047] Typically, the dosage range of chromium administered to an individual in the form of chromium picolinate, chromium nicotinate, or other chromium complex provides between about 50 and 10,000 micrograms per day of chromium; preferably between about 100 and 1,000 micrograms per day; more preferably, between about 200 and 500 micrograms per day.

[0048] In some embodiments, methods of increasing glucose uptake by cells or by organisms, such as mammals and humans, treating insulin-dependent diabetes, reducing body fat, improving insulin sensitivity, reducing hyperglycemia, and reducing hypercholesterolemia with chromium complexes and conjugated compounds such as conjugated fatty acids or conjugated fatty alcohols are contemplated and isomers of fatty acids, as well as isomers of fatty alcohols. Advantageously, a subject is administered a pharmaceutically effective dose of a chromium complex. In one embodiment, the conjugated compound is administered substantially simultaneously with the chromium complex. In an alternative embodiment, the chromium complex is administered first and then the conjugated compound is added second. In yet another embodiment, the conjugated compound is administered first. Preferably, the chromium complex and conjugated compound are administered to a subject within twenty-four hours of each other. In a particularly preferred embodiment, the chromium complex and conjugated compound are administered within an hour of each other.

[0049] The following examples teach certain methods and compositions for treating diabetes and obesity through the administration of at least one chromium complex in concert with a conjugated compound. The subject can advantageously be a vertebrate, a mammal, a bird, and is preferably a human, a cat, or a dog. It will be appreciated, however, that the at least one chromium complex may be administered first, then the conjugated compound. Alternatively, the conjugated compound can be administered first, then the chromium complex. As illustrated in the following examples, the composition may optionally include picolinic acid, nicotinic acid, or both picolinic acid and nicotinic acid in combination with a chromium complex and conjugated compound. These examples are illustrative only and are not intended to limit the scope of the invention disclosed herein. The treatment methods described below can be optimized using empirical techniques well known to those of ordinary skill in the art. Moreover, artisans of skill would be able to use the teachings described in the following examples to practice the full scope of the invention disclosed herein.

#### EXAMPLE 1

[0050] An obese adult human subject is identified. The subject is orally administered a tablet containing about 300 µg chromium as chromium picolinate and 1 gram conjugated linoleic acid twice a day. The tablet additionally comprises

ibuprofen in a pharmaceutically effective dose of 200 mg. Over the course of several weeks, a decrease in body mass is observed. The chromium picolinate in combination with conjugated linoleic acid synergistically reduce the subject's body mass.

[0051] It will be appreciated that although specific embodiments of the invention have been described herein for the purposes of illustration, various modifications may be made without deviating from the spirit and scope of the invention. Accordingly, the invention is not limited except as by the appended claims.

## EXAMPLE 2

[0052] A subject suffering from insulin-dependent diabetes is identified. The subject is orally administered a daily dose of one tablet containing about 500 µg chromium as chromium nicotinate and 500 mg conjugated linoleic acid. Over the course of several days, an improvement in glucose uptake in the subject is observed and insulin dependence is reduced. The chromium nicotinate in combination with linoleic acid act synergistically to improve the subject's glucose tolerance and to treat the subjects diabetes.

## Claims

1. A composition for reducing body fat, improving insulin sensitivity, reducing hyperglycemia, and reducing hypercholesterolemia, said composition consisting essentially of at least one chromium complex and a conjugated fatty acid or conjugated fatty alcohol.
2. The composition of Claim 1, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrienoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.
3. The composition of Claim 1, wherein said conjugated fatty acid is conjugated linoleic acid.
4. The composition of Claim 1, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, docosapentaenoic alcohol, and all other diunsaturated and polyunsaturated fatty alcohols.
5. The composition of claim 1, wherein said chromium complex is chromium picolinate or chromium nicotinate.
6. A composition for reducing body fat, improving insulin sensitivity, reducing hyperglycemia, and reducing hypercholesterolemia, said composition consisting essentially of at least one chromium complex, a conjugated fatty acid or conjugated fatty alcohol, and a chelating agent.
7. The composition of claim 6, wherein said chelating agent is picolinic acid, nicotinic add, or both.
8. The composition of claim 1, wherein said composition is incorporated into a pharmaceutically effective carrier.
9. The composition of claim 8, wherein said pharmaceutically effective carrier is selected from the group consisting of a tablet, capsule, microbead, emulsion, powder, granule, suspension, syrup and elixir.
10. The composition of claim 9, wherein said microbead is a sugar beadlet or microcrystalline cellulose beadlet and said at least one chromium complex and said conjugated linoleic acid are coated on said beadlet.
11. The composition of claim 9, wherein said tablet, capsule, or microbead is coated with an enteric coating.
12. The composition of claim 1, wherein said chromium complex and said conjugated fatty acid or conjugated fatty alcohol are in a ratio of between about 1:3000 to 1:50 (w/w).

13. A composition for reducing body fat, improving insulin sensitivity, reducing hyperglycemia, and reducing hypercholesterolemia, said composition consisting essentially of at least one chromium complex, a conjugated fatty acid or conjugated fatty alcohol and at least one of a cyclooxygenase inhibitor, a mucolytic, and a salicin-containing herb.

14. The composition of claim 13, wherein said cyclooxygenase inhibitor is selected from the group consisting of indomethacin, ibuprofen, acetaminophen, and naproxen.

15. The composition of claim 13, wherein said salicin-containing herb is selected from the group consisting of *Boswellia serrata* (frankincense), *Betula lenta* (sweet birch), *Betula pubescens* (white birch), *Filipendula ulmaria* (meadowsweet), *Gautheria procumbens* (wintergreens), *Populus balsamifera*, *Populus jackii* (balm of Gilead) and *Salix alba* (white willow).

16. The composition of claim 13, wherein said mucolytic is guaifenesin.

17. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts and a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol in the manufacture of a medicament for administration to a subject in need thereof for the treatment of obesity.

18. The use of claim 17, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrenoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.

19. The use of claim 17, wherein said conjugated fatty acid is conjugated linoleic acid.

20. The use of claim 17, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrenoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, docosapentaenoic alcohol, and all other diunsaturated and polyunsaturated fatty alcohols.

21. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts, a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol, and at least one uncomplexed chelating agent in the manufacture of a medicament for administration to a subject in need thereof for the treatment of obesity.

22. The use of claim 21, wherein said chelating agent is picolinic acid, nicotinic acid, or both.

23. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts and a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol in the manufacture of a medicament for administration to a subject in need thereof for reducing body fat.

24. The use of claim 23, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrenoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.

25. The use of claim 23, wherein said conjugated fatty acid is conjugated linoleic acid.

26. The use of claim 23, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated

version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, docosapentaenoic alcohol, and all other diunsaturated and polyunsaturated fatty alcohols.

27. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts, a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol, and at least one uncomplexed chelating agent in the manufacture of a medicament for administration to a subject in need thereof for reducing body fat.

28. The use of claim 27, wherein said chelating agent is picolinic acid, nicotinic acid, or both.

29. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts and a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol in the manufacture of a medicament for administration to a subject in need thereof for improving insulin sensitivity.

30. The use of claim 29, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrienoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.

31. The use of claim 29, wherein said conjugated fatty acid is conjugated linoleic acid.

32. The use of claim 29, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, and docosapentaenoic alcohol.

33. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts, a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol, and at least one uncomplexed chelating agent in the manufacture of a medicament for administration to a subject in need thereof for improving insulin sensitivity.

34. The use of claim 33, wherein said chelating agent is picolinic acid, nicotinic acid, or both.

35. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts and a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol in the manufacture of a medicament for administration to a subject in need thereof for reducing hyperglycemia.

36. The use of claim 35, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrienoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.

37. The use of claim 35, wherein said conjugated fatty acid is conjugated linoleic acid.

38. The use of claim 35, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatetraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, and docosapentaenoic alcohol.

donic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, and docosapentaenoic alcohol.

39. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts, a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol, and at least one uncomplexed chelating agent in the manufacture of a medicament for administration to a subject in need thereof for reducing hyperglycemia.

40. The use of claim 39, wherein said chelating agent is picolinic acid, nicotinic acid, or both.

41. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts and a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol in the manufacture of a medicament for administration to a subject in need thereof for reducing hypercholesterolemia.

42. The use of claim 41, wherein said conjugated fatty acid is selected from the group consisting of a conjugated version of linoleic acid, linolenic acid, gamma linolenic acid, arachidonic acid, mead acid, stearidonic acid, alpha-eleostearic acid, eleostearic acid, pinolenic acid, docosadienic acid, docosatetraenoic acid, octadecadienoic acid, octadecatrienoic acid, eicosatetraenoic acid, eicosapentaenoic acid, docosahexaenoic acid, and docosapentaenoic acid.

43. The use of claim 41, wherein said conjugated fatty acid is conjugated linoleic acid.

44. The use of claim 41, wherein said conjugated fatty alcohol is selected from the group consisting of a conjugated version of linoleic alcohol, linolenic alcohol, gamma linolenic alcohol, arachidonic alcohol, mead alcohol, stearidonic alcohol, alpha-eleostearic alcohol, eleostearic alcohol, pinolenic alcohol, docosadienic alcohol, docosatraenoic alcohol, octadecadienoic alcohol, octadecatrienoic alcohol, eicosatetraenoic alcohol, eicosapentaenoic alcohol, docosahexaenoic alcohol, and docosapentaenoic alcohol.

45. Use of a composition consisting essentially of a pharmaceutically effective dose of chromium complex selected from the group consisting of chromium picolinate, chromium nicotinate, chromic tripicolinate, chromic polynicotinate, chromium chloride, chromium histidinate, and chromium yeasts, a pharmaceutically effective dose of conjugated fatty acid or conjugated fatty alcohol, and at least one uncomplexed chelating agent in the manufacture of a medicament for administration to a subject in need thereof for reducing hypercholesterolemia.

46. The use of claim 45, wherein said chelating agent is picolinic acid, nicotinic acid, or both.

## Revendications

1. Composition pour réduire le tissu adipeux, améliorer la sensibilité à l'insuline, réduire l'hyperglycémie et réduire l'hypercholestérolémie, ladite composition consistant essentiellement en au moins un complexe de chrome et un acide gras conjugué ou alcool gras conjugué.

2. Composition suivant la revendication 1, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linolénique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétranoïque, d'acide octadécadiénique, d'acide octadécatriénique, d'acide eicosatétranoïque, d'acide eicosapentaénique, d'acide docosahexaénique, et d'acide docosapentaénique.

3. Composition suivant la revendication 1, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.

4. Composition suivant la revendication 1, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linolénique, d'alcool gamma-linolénique, d'alcool arachi-

donique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétranoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétraénoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque, d'alcool docosapentaénoïque et de tous les autres alcools gras diinsaturés et polyinsaturés.

- 5 5. Composition suivant la revendication 1, dans laquelle ledit complexe de chrome est le picolinate de chrome ou le nicotinate de chrome.
- 10 6. Composition pour réduire le tissu adipeux, améliorer la sensibilité à l'insuline, réduire l'hyperglycémie et réduire l'hypercholestérolémie, ladite composition consistant essentiellement en au moins un complexe de chrome, un acide gras conjugué ou un alcool gras conjugué, et un agent chélatant.
- 15 7. Composition suivant la revendication 6, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique ou ces deux acides.
- 20 8. Composition suivant la revendication 1, ladite composition étant incorporée à un support pharmaceutiquement efficace.
9. Composition suivant la revendication 8, dans laquelle ledit support pharmaceutiquement efficace est choisi dans le groupe consistant en un comprimé, une capsule, une microbille, une émulsion, une poudre, un granule, une suspension, un sirop et un élixir.
- 25 10. Composition suivant la revendication 9, dans laquelle ladite microbille est une perle de sucre, ou une perle de cellulose microcristalline, et ledit au moins un complexe de chrome et ledit acide linoléique conjugué sont présents sous forme de revêtement sur ladite perle.
11. Composition suivant la revendication 9, dans laquelle ledit comprimé, ladite capsule ou ladite microbille est revêtue d'un enrobage entérique.
- 30 12. Composition suivant la revendication 1, dans laquelle ledit complexe de chrome et ledit acide gras conjugué ou alcool gras conjugué sont présents en un rapport compris dans l'intervalle d'environ 1:3000 à 1:50 (poids/poids).
- 35 13. Composition pour réduire le tissu adipeux, améliorer la sensibilité à l'insuline, réduire l'hyperglycémie et réduire l'hypercholestérolémie, ladite composition consistant essentiellement en au moins un complexe de chrome, un acide gras conjugué ou alcool gras conjugué et au moins un des agents consistant en un inhibiteur de cyclooxygénase, un agent mucolytique et une plante herbacée contenant de la salicine.
- 40 14. Composition suivant la revendication 13, dans laquelle ledit inhibiteur de cyclooxygénase est choisi dans le groupe consistant en l'indométhacine, l'ibuprofène, l'acétaminophène et le naproxène.
- 45 15. Composition suivant la revendication 13, dans laquelle ladite plante herbacée contenant de la salicine est choisie dans le groupe consistant en *Boswellia serrata* (encens), *Betula lenta* (bouleau à sucre), *Betula pubescens* (bouleau pubescent), *Filipendula ulmaria* (reine des prés), *Gautheria procumbens* (gaulthérie), *Populus balsamifera*, *Populus jackii* (baume de Gilead) et *Salix alba* (saule blanc).
16. Composition suivant la revendication 13, dans laquelle ledit agent mucolytique et la guaifénésine.
- 50 17. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué dans la production d'un médicament destiné à l'administration à un sujet nécessitant cette administration pour le traitement de l'obésité.
- 55 18. Utilisation suivant la revendication 17, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linolénique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétranoïque, d'acide octadécadiénoïque, d'acide octadécatriénoïque, d'acide eicosatétraénoïque, d'acide eicosapentaénoïque, d'acide docosahexaénoïque et d'acide do-

cosapentaénoïque.

19. Utilisation suivant la revendication 17, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.

20. Utilisation suivant la revendication 17, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linoléinique, d'alcool gamma-linolénique, d'alcool arachidonique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétranoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétranoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque, d'alcool docosapentaénoïque, et de tous les autres alcools gras diinsaturés et polyinsaturés.

21. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome, et levures au chrome, et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué, et au moins un agent chélatant non complexé dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour le traitement de l'obésité.

22. Utilisation suivant la revendication 21, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique ou ces deux acides.

23. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour réduire le tissu adipeux.

24. Utilisation suivant la revendication 23, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linoléinique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétranoïque, d'acide octadécadiénoïque, d'acide octadécatriénoïque, d'acide eicosatétranoïque, d'acide eicosapentaénoïque, d'acide docosahexaénoïque et d'acide docosapentaénoïque.

25. Utilisation suivant la revendication 23, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.

26. Utilisation suivant la revendication 23, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linoléinique, d'alcool gamma-linolénique, d'alcool arachidonique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétranoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétranoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque, d'alcool docosapentaénoïque, et de tous les autres alcools gras diinsaturés et polyinsaturés.

27. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome, et levures au chrome, une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué, et au moins un agent chélatant non complexé dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour la réduction du tissu adipeux.

28. Utilisation suivant la revendication 27, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique ou ces deux acides.

29. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour améliorer la sensibilité à l'insuline.

30. Utilisation suivant la revendication 29, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linolénique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétranoïque, d'acide octadécadiénoïque, d'acide octadécatriénoïque, d'acide eicosatétraénoïque, d'acide eicosapentaénoïque, d'acide docosahexaénoïque et d'acide docosapentaénoïque.

31. Utilisation suivant la revendication 29, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.

32. Utilisation suivant la revendication 29, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linolénique, d'alcool gamma-linolénique, d'alcool arachidonique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétranoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétraénoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque et d'alcool docosapentaénoïque.

33. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué, et au moins un agent chélatant non complexé dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour améliorer la sensibilité à l'insuline.

34. Utilisation suivant la revendication 33, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique, ou ces deux acides.

35. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour réduire l'hyperglycémie.

36. Utilisation suivant la revendication 35, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linolénique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétranoïque, d'acide octadécadiénoïque, d'acide octadécatriénoïque, d'acide eicosatétraénoïque, d'acide eicosapentaénoïque, d'acide docosahexaénoïque et d'acide docosapentaénoïque.

37. Utilisation suivant la revendication 35, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.

38. Utilisation suivant la revendication 35, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linolénique, d'alcool gamma-linolénique, d'alcool arachidonique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétranoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétraénoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque et d'alcool docosapentaénoïque.

39. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome, et levures au chrome, une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué, et au moins un agent chélatant non complexé dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour réduire l'hyperglycémie.

40. Utilisation suivant la revendication 39, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique ou ces deux acides.



41. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome et une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour réduire l'hypercholestérolémie.
42. Utilisation suivant la revendication 41, dans laquelle ledit acide gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'acide linoléique, d'acide linoléique, d'acide gamma-linolénique, d'acide arachidonique, d'acide d'hydromel, d'acide stéaridonique, d'acide alpha-éléostéarique, d'acide éléostéarique, d'acide pinolénique, d'acide docosadiénique, d'acide docosatétraténoïque, d'acide octadécadiénoïque, d'acide octadécatriénoïque, d'acide eicosatétraénoïque, d'acide eicosapentaénoïque, d'acide docosahexaénoïque et d'acide docosapentaénoïque.
43. Utilisation suivant la revendication 41, dans laquelle ledit acide gras conjugué est l'acide linoléique conjugué.
44. Utilisation suivant la revendication 41, dans laquelle ledit alcool gras conjugué est choisi dans le groupe consistant en des versions conjuguées d'alcool linoléique, d'alcool linoléique, d'alcool gamma-linolénique, d'alcool arachidonique, d'alcool d'hydromel, d'alcool stéaridonique, d'alcool alpha-éléostéarique, d'alcool éléostéarique, d'alcool pinolénique, d'alcool docosadiénique, d'alcool docosatétraténoïque, d'alcool octadécadiénoïque, d'alcool octadécatriénoïque, d'alcool eicosatétraénoïque, d'alcool eicosapentaénoïque, d'alcool docosahexaénoïque et d'alcool docosapentaénoïque.
45. Utilisation d'une composition consistant essentiellement en une dose pharmaceutiquement efficace d'un complexe de chrome choisi dans le groupe consistant en picolinate de chrome, nicotinate de chrome, tripicolinate chromique, polynicotinate chromique, chlorure de chrome, histidinate de chrome et levures au chrome, une dose pharmaceutiquement efficace d'un acide gras conjugué ou alcool gras conjugué, et au moins un agent chélatant non complexé dans la production d'un médicament destiné à l'administration à un sujet nécessitant une telle administration pour réduire l'hypercholestérolémie.
46. Utilisation suivant la revendication 45, dans laquelle ledit agent chélatant est l'acide picolinique, l'acide nicotinique ou ces deux acides.

#### Patentansprüche

1. Zusammensetzung zur Verminderung des Körperfettes, zur Verbesserung der Insulinempfindlichkeit, zur Verminderung von Hyperglykämie und zur Verminderung von Hypercholesterolämie, wobei diese Zusammensetzung im wesentlichen aus einem Chromkomplex und einer konjugierten Fettsäure oder einem konjugierten Fettalkohol besteht.
2. Zusammensetzung nach Anspruch 1, worin die konjugierte Fettsäure aus der Gruppe ausgewählt ist, die aus einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Melsäure, Stearidonsäure, alpha-Elaeostearinsäure, Elaeostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadeciadiensäure, Octadecatransäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure besteht.
3. Zusammensetzung nach Anspruch 1, worin die konjugierte Fettsäure konjugierte Linolsäure ist.
4. Zusammensetzung nach Anspruch 1, worin der konjugierte Fettalkohol aus der Gruppe ausgewählt ist, die aus einer konjugierten Version von Linolalkohol, Linolenalkohol, gamma-Linolenalkohol, Arachidonalkohol, Metalkohol, Stearidonalkohol, alpha-Elaeostearinalkohol, Elaeostearinalkohol, Pinolenalkohol, Docosadienalkohol, Docosatetraenalkohol, Octadecadienalkohol, Octadecatrienalkohol, Eicosatetraenalkohol, Eicosapentaenalkohol, Docosahexaenalkohol, Docosapentaenalkohol und allen anderen di-ungesättigten und polyungesättigten Fettalkoholen besteht.
5. Zusammensetzung nach Anspruch 1, worin der Chromkomplex Chrompicolinat oder Chromnicotinat ist.
6. Zusammensetzung zum Vermindern des Körperfettes, zur Verbesserung der Insulinempfindlichkeit, Verminderung

der Hyperglykämie und Verminderung der Hypercholesterolämie, wobei diese Zusammensetzung im wesentlichen aus wenigstens einem Chromkomplex, einer konjugierten Fettsäure oder einem konjugierten Fettalkohol sowie einem Chelatisierungsmittel besteht.

- 5 7. Zusammensetzung nach Anspruch 6, worin das Chelatisierungsmittel Picolinsäure, Nicotinsäure oder beides ist.
8. Zusammensetzung nach Anspruch 1, worin die Zusammensetzung in einen pharmazeutisch wirksamen Träger eingearbeitet ist.
- 10 9. Zusammensetzung nach Anspruch 8, worin der pharmazeutisch wirksame Träger aus der Gruppe ausgewählt ist, die aus einer Tablette, einer Kapsel, einer Mikroperle, einer Emulsion, einem Pulver, einem Granulat, einer Suspension, einem Sirup und einem Elixier besteht.
- 15 10. Zusammensetzung nach Anspruch 9, worin die Mikroperle ein Zuckerperlchen oder mikrokristallines Celluloseperlchen ist und wenigstens ein Chromkomplex und die konjugierte Linolsäure als Überzug auf dem Perlchen sich befinden.
11. Zusammensetzung nach Anspruch 9, worin die Tablette, Kapsel oder Mikroperle mit einem enteralem Überzug versehen ist.
- 20 12. Zusammensetzung nach Anspruch 1, worin der Chromkomplex und die konjugierte Fettsäure oder der konjugierte Fettalkohol in einem Verhältnis zwischen etwa 1 : 3000 und 1 : 50 (Gewicht/Gewicht) vorliegen.
- 25 13. Zusammensetzung zur Verminderung des Körperfettes, Verbesserung der Insulinempfindlichkeit, Verminderung der Hyperglykämie und Verminderung der Hypercholesterolämie, wobei diese Zusammensetzung im wesentlichen aus wenigstens einem Chromkomplex, einer konjugierten Fettsäure oder einem konjugierten Fettalkohol sowie wenigstens einem Cyclooxygenaseinhibitor, einem Schleimlöser und einer salicinhaltigen Pflanze besteht.
- 30 14. Zusammensetzung nach Anspruch 13, worin der Cyclooxygenaseinhibitor aus der Gruppe ausgewählt ist, die aus Indometacin, Ibuprofen, Acetaminophen und Naproxen besteht.
- 35 15. Zusammensetzung nach Anspruch 13, worin die salicinhaltige Pflanze aus der Gruppe ausgewählt ist, die aus *Boswellia serrata* (Weihrauchbaum), *Betula lenta* (Zuckerbirke), *Betula pubescens* (Moorbirke), *Filipendula ulmaria* (Echtes Mädesüß), *Gaultheria procumbens* (Scheinbeere), *Populus balsamifera* (Balsampappel), *Populus jackii* (Sarg) und *Salix alba* (Silberweide) besteht.
16. Zusammensetzung nach Anspruch 13, worin der Schleimlöser Guafenesin ist.
- 40 17. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe sowie einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol bei der Herstellung eines Arzneimittels für Verabreichung an einen Patienten, der dieses für die Behandlung von Fettleibigkeit braucht.
- 45 18. Verwendung nach Anspruch 17, bei der die konjugierte Fettsäure aus der Gruppe ausgewählt ist, die aus einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Methylsäure, Stearidonsäure, alpha-Elaeostearinsäure, Elaeostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadecadiensäure, Octadecatriensäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure besteht.
- 50 19. Verwendung nach Anspruch 17, worin die konjugierte Fettsäure konjugierte Linolsäure ist.
- 55 20. Verwendung nach Anspruch 17, worin der konjugierte Fettalkohol aus der Gruppe ausgewählt ist, die aus der konjugierten Version von Linolalkohol, Linotenalkohol, gamma-Linolenalkohol, Arachidonalkohol, Metalkohol, Stearidonalkohol, alpha-Elaeostearinalkohol, Elaeostearinalkohol, Pinolenalkohol, Docosadienalkohol, Docosatetraenalkohol, Octadecadienalkohol, Octadecatrienalkohol, Eicosatetraenalkohol, Eicosapentaenalkohol, Docosahexaenalkohol, Docosapentaenalkohol und allen anderen di-ungesättigten und polyungesättigten Fettalkoholen besteht.

21. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe ausgewählt ist, die aus Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrombrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe besteht, einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder von konjugiertem Fettalkohol und wenigstens einem unkomplexierten Chelatisiermittel besteht, bei der Herstellung eines Arzneimittels zur Verabreichung an einen Patienten, der dieses für die Behandlung von Fettleibigkeit braucht.
22. Verwendung nach Anspruch 21, wobei das Chelatisiermittel Picolinsäure, Nicotinsäure oder beides ist.
23. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe ausgewählt ist, die aus Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe besteht, und einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol bei der Herstellung eines Arzneimittels für die Verabreichung an einen Patienten, der dieses zur Verminderung von Körperfett braucht.
24. Verwendung nach Anspruch 23, bei der die konjugierte Fettsäure aus der Gruppe ausgewählt ist, die aus einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Melsäure, Stearidonsäure, alpha-Eicostearinsäure, Eicostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadecadiensäure, Octadecatriensäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure besteht.
25. Verwendung nach Anspruch 23, wobei die konjugierte Fettsäure konjugierte Linolsäure ist.
26. Verwendung nach Anspruch 23, bei der der konjugierte Fettalkohol aus der Gruppe ausgewählt ist, die aus einer konjugierten Version von Linolalkohol, Linolenalkohol, gamma-Linolenalkohol, Arachidonalkohol, Metalkohol, Stearidonalkohol, alpha-Elaeostearinalkohol, Elaeostearinalkohol, Pinolenalkohol, Docosadienalkohol, Docosatetraenalkohol, Octadecadienalkohol, Octadecatrienalkohol, Eicosatetraenalkohol, Eicosapentaenalkohol, Docosahexaenalkohol, Docosapentaenalkohol und allen anderen di-ungesättigten und polyungesättigten Fettsäuren besteht.
27. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe ausgewählt ist, die aus Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe besteht, einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol und wenigstens einem unkomplexierten Chelatisiermittel besteht, bei der Herstellung eines Arzneimittels für die Verabreichung an einen Patienten, der dieses braucht zur Verminderung von Körperfett.
28. Verwendung nach Anspruch 27, bei der das Chelatisiermittel Picolinsäure, Nicotinsäure oder beides ist.
29. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe ausgewählt ist, und einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol bei der Herstellung eines Arzneimittels zur Verabreichung an einen Patienten, der es zur Verbesserung der Insulinempfindlichkeit benötigt.
30. Verwendung nach Anspruch 29, bei der die konjugierte Fettsäure aus der Gruppe einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Melsäure, Stearidonsäure, alpha-Elaeostearinsäure, Elaeostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadecadiensäure, Octadecatriensäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure ausgewählt ist.
31. Verwendung nach Anspruch 29, bei der die konjugierte Fettsäure konjugierte Linolsäure ist.
32. Verwendung nach Anspruch 29, bei der der konjugierte Fettalkohol aus der Gruppe einer konjugierten Version von Linolalkohol, Linolenalkohol, gamma-Linolenalkohol, Arachidonalkohol, Metalkohol, Stearidonalkohol, alpha-Elaeostearinalkohol, Elaeostearinalkohol, Pinolenalkohol, Docosadienalkohol, Docosatetraenalkohol, Octadecadienalkohol, Octadecatrienalkohol, Eicosatetraenalkohol, Eicosapentaenalkohol, Docosahexaenalkohol und Docosapentaenalkohol ausgewählt ist.

33. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe ausgewählt ist, einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol und wenigstens einem unkomplexierten Chelatierungsmittel besteht, bei der Herstellung eines Arzneimittels für die Verabreichung an einen Patienten, der es zur Verbesserung der Insulinempfindlichkeit braucht.
34. Verwendung nach Anspruch 33, bei der das Chelatierungsmittel Picolinsäure, Nicotinsäure oder beides ist.
35. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe und einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol besteht, bei der Herstellung eines Arzneimittels für die Verabreichung an einen Patienten, der es für die Verminderung von Hyperglykämie benötigt.
36. Verwendung nach Anspruch 35, bei der die konjugierte Fettsäure aus der Gruppe einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Methylsäure, Stearidonsäure, alpha-Elaeostearinsäure, Elaeostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadecadiensäure, Octadecatriensäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure besteht.
37. Verwendung nach Anspruch 35, worin die konjugierte Fettsäure konjugierte Linolsäure ist.
38. Verwendung nach Anspruch 35, bei welcher der konjugierte Fettalkohol aus der Gruppe einer konjugierten Version von Linolalkohol, Linolenalkohol, gamma-Linolenalkohol, Arachidonalkohol, Metalkohol, Stearidonalkohol, alpha-Elaeostearinalkohl, Elaeostearinalkohl, Pinolenalkohl, Docosadienalkohl, Docosatetraenalkohl, Octadecadienalkohl, Octadecatrienalkohl, Eicosatetraenalkohl, Eicosapentaenalkohl, Docosahexaenalkohl und Docosapentaenalkohl ausgewählt ist.
39. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe ausgewählt ist, und einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol und wenigstens einem unkomplexierten Chelatierungsmittel besteht, bei der Herstellung eines Arzneimittels für die Verabreichung an einen Patienten, der es für die Verminderung von Hyperglykämie benötigt.
40. Verwendung nach Anspruch 39, bei der das Chelatierungsmittel Picolinsäure, Nicotinsäure oder beides ist.
41. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex, der aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chromchlorid, Chromhistidinat und Chromhefe ausgewählt ist, und einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohl besteht, bei der Herstellung eines Arzneimittels zur Verabreichung an einen Patienten, der es zur Verminderung von Hypercholesterolemie benötigt.
42. Verwendung nach Anspruch 41, bei der die konjugierte Fettsäure aus der Gruppe einer konjugierten Version von Linolsäure, Linolensäure, gamma-Linolensäure, Arachidonsäure, Methylsäure, Stearidonsäure, alpha-Elaeostearinsäure, Elaeostearinsäure, Pinolensäure, Docosadiensäure, Docosatetraensäure, Octadecadiensäure, Octadecatriensäure, Eicosatetraensäure, Eicosapentaensäure, Docosahexaensäure und Docosapentaensäure besteht.
43. Verwendung nach Anspruch 41, bei der die konjugierte Fettsäure konjugierte Linolsäure ist.
44. Verwendung nach Anspruch 41, bei der die konjugierte Fettsäure aus der Gruppe ausgewählt ist, die eine konjugierte Version von Linolalkohl, Linolenalkohl, gamma-Linolenalkohl, Arachidonalkohl, Metalkohl, Stearidonalkohl, alpha-Elaeostearinalkohl, Elaeostearinalkohl, Pinolenalkohl, Docosadienalkohl, Docosatetraenalkohl, Octadecadienalkohl, Octadecatrienalkohl, Eicosatetraenalkohl, Eicosapentaenalkohl, Docosahexaenalkohl und Docosapentaenalkohl besteht.
45. Verwendung einer Zusammensetzung, die im wesentlichen aus einer pharmazeutisch wirksamen Dosis von Chromkomplex aus der Gruppe Chrompicolinat, Chromnicotinat, Chromtripicolinat, Chrompolynicotinat, Chrom-

## EP 1 357 977 B1

chlorid, Chromhistidinat und Chromhefe ausgewählt ist, einer pharmazeutisch wirksamen Dosis von konjugierter Fettsäure oder konjugiertem Fettalkohol und wenigstens einem unkomplexierten Chelatisiermittel besteht, bei der Herstellung eines Arzneimittels zur Verabreichung an einen Patienten, der es zur Verminderung seiner Hypercholesterolemie benötigt

5

46. Verwendung nach Anspruch 45, bei der das Chelatisiermittel Picolinsäure, Nicotinsäure oder beides ist.

10

15

20

25

30

35

40

45

50

55

## The Effect of Conjugated Linoleic Acid (CLA) on Glucose Uptake in Human Skeletal Muscle Culture

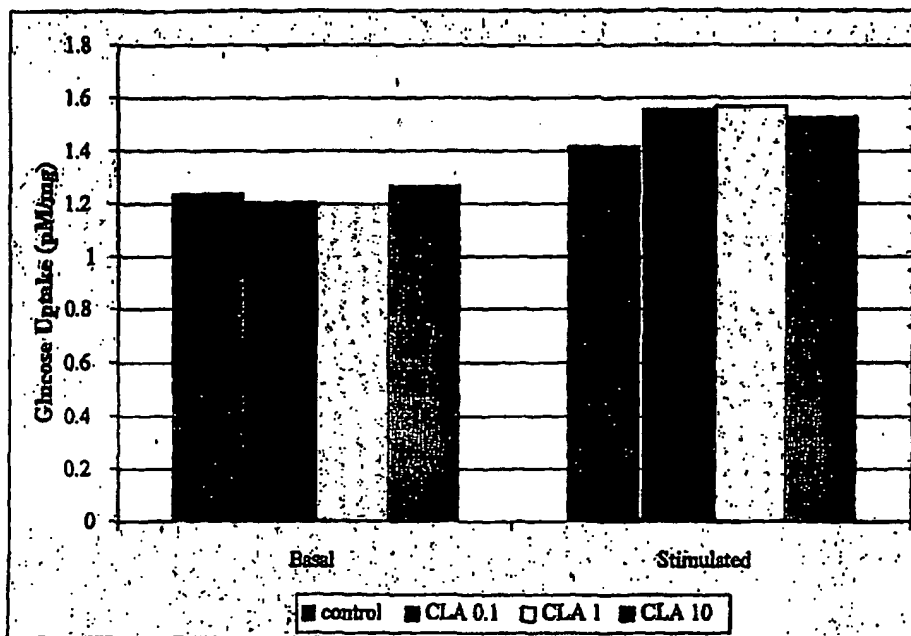
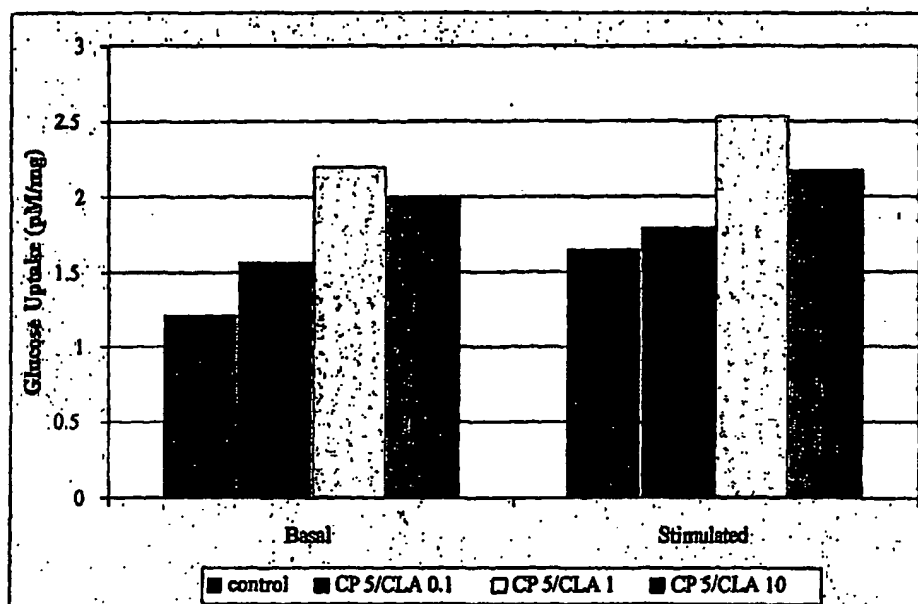


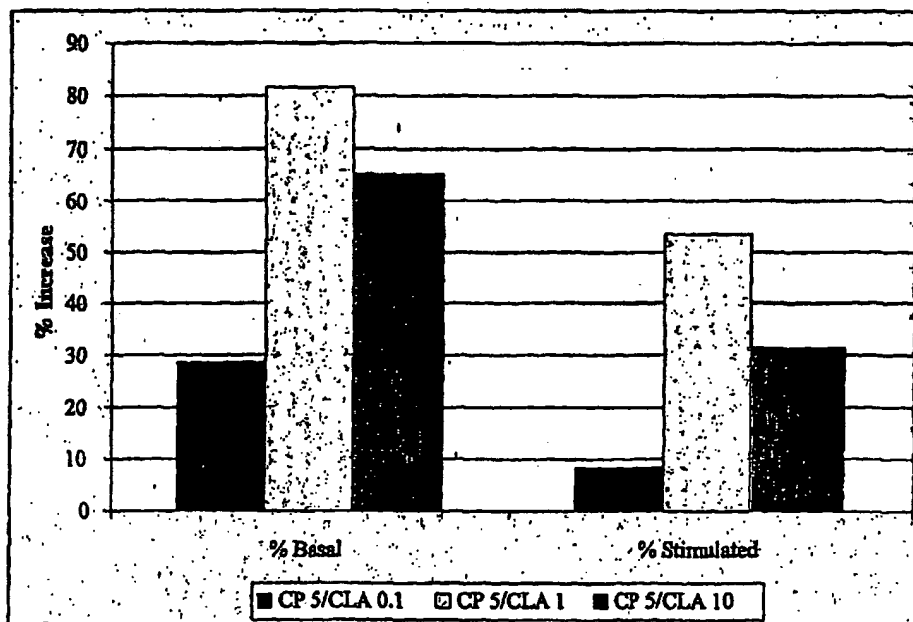
FIGURE 1

## The Effect of CP+CLA on Glucose Uptake in Human Skeletal Muscle Culture



**FIGURE 2**

# **The Effect of CP+CLA on Glucose Uptake in Human Skeletal Muscle Culture** (Percent Increase Over Baseline)



**FIGURE 3**